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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE  
Group Art Unit 1614

In re

Patent Application of

David Edwin Thurston, et al.

Application No. 10/534,825 ✓

Confirmation No.: 8880

Filed: October 20, 2005

Examiner: Unknown

"PYRROLOBENZODIAZEPINES"

SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT  
PURSUANT TO 37 CFR §1.97(b)

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Sir:

The Examiner's attention is directed to the references which are listed on the attached Form PTO/SB/08A and PTO/SB/08B and copies of non-U.S. patent references are attached.

Citation of these references is respectfully requested.

No concession is made that these documents are prior art, and Applicant expressly reserves the right to antedate the documents as may be appropriate.

Respectfully submitted,

*Charlene L. Yager*  
Charlene L. Yager  
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File No. 065435-9048-US00

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Substitute for form 1449/PTO <b>INFORMATION DISCLOSURE STATEMENT BY APPLICANT</b> APR 26 2006 (use as many sheets as necessary)		<b>Complete if Known</b>	
		Application Number	10/534,825
		Filing Date	October 20, 2005
		First named Inventor	David Edwin Thurston
		Group Art Unit	1614
		Examiner name	Unknown
		Attorney Docket Number	065435-9048-US00

**U.S. Patent Documents**

Examiner Initials		U.S. Patent Document Number	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document
		6,660,856	Wang	12/9/2003

**FOREIGN PATENT DOCUMENTS**

Examiner Initials	Country Code	Foreign Patent Document Number	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document	Translation	English Abstract
	EP	1193270	Spirogen Ltd.	4/3/2002		
	GB	2053894	The Green Cross Corporation	2/11/1981		
	WO	88/04659	The Upjohn Company	6/30/1988		
	WO	91/16324	The Upjohn Company	10/31/1991		
	WO	96/23497	Synphar Laboratories, Inc.	8/8/1996		
	WO	97/07097	Auckland Division Cancer Society of New Zealand Inc.	2/27/1997		
	WO	98/11101	Auckland Division Cancer Society of New Zealand Inc.	3/19/1998		
	WO	98/12197	Kyorin Pharmaceuticals Co., Ltd. et al.	3/26/1998	N	Y
	WO	99/29642	The Scripps Research Institute	6/17/1999		
	WO	99/46244	Novo Nordisk A/S et al.	9/16/1999		

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(use as many sheets as necessary)		<b>Filing Date</b>	October 20, 2005
Sheet 2 of 2		<b>First named Inventor</b>	David Edwin Thurston
		<b>Group Art Unit</b>	1614
		<b>Examiner name</b>	Unknown
		<b>Attorney Docket Number</b>	065435-9048-US00

**U.S. Patent Documents****FOREIGN PATENT DOCUMENTS**

Examiner Initials		Country Code	Foreign Patent Document Number	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document	Translation	English Abstract
		WO	00/12508	The University of Portsmouth Higher Education Corp.	3/9/2000		
		WO	00/12509	The University of Portsmouth Higher Education Corp.	3/9/2000		
		WO	00/64864	Cancer Campaign Research Technology Ltd.	11/2/2000		
		WO	2004/043963	Spirogen Ltd.	5/27/2004		
		WO	2005/023814	Spirogen Ltd.	3/17/2005		
		WO	2005/040170	Government of the U.S.A. et al.	5/6/2005		
		WO	2005/085251	Spirogen Ltd.	9/15/2005		

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Information Disclosure Statement for form 1449B/PTO**INFORMATION DISCLOSURE  
STATEMENT BY APPLICANT**

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Sheet	1	of	5	Attorney Docket Number	065435-9048-US00
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**OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS**

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		ADAMS et al., "Molecular modelling of a sequence-specific DNA-binding agent based on the pyrrolo[2,1-c][1,4]benzodiazepines," Pharm. Pharmacol. Commun. (1999) 5:555-560
		BARALDI, P.G. et al., "[2,1-c][1,4]benzodiazepine (PBD)-distamycin hybrid inhibits DNA binding to transcription factor Sp1," Nucleotides and Nucleic Acids (2000) 19(8):1219-1229
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		DE GROOT, FMH et al., "Synthesis and biological evaluation of 2'-carbamate-linked 2'-carbonate-linked prodrugs of paclitaxel: selective activation by the tumor-associated protease plasmin," J. Med. Chem. (2000) 43(16):3093-3102
		DE GROOT, FMH et al., "Novel 20-carbonate linked prodrugs of camptothecin and 9-aminocamptothecin designed for activation by tumour-associated plasmin," Biorg. Med. Chem. Lett. (2002) 12(17):2371-2376

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				First Named Inventor	David Edwin Thurston
				Group Art Unit	1614
				Examiner Name	Unknown
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		DUBOWCHIK, G.M. et al., "Cathepsin B-sensitive dipeptide prodrugs. 1. A model study of structural requirements for efficient release of doxorubicin," <i>Biorg. Med. Chem. Lett.</i> (1998) 8:3341-3346
		DUBOWCHIK, G.M. et al., "Cathepsin B-sensitive dipeptide prodrugs. 2. Models of anticancer drugs paclitaxel (Taxol), Mitomycin C and Doxorubicin," <i>Biorg. Med. Chem. Lett.</i> (1998) 8:3347-3352
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		GARSKY et al., "The synthesis of a prodrug of doxorubicin designed to provide reduced systemic toxicity and greater target efficacy," <i>J. Med. Chem.</i> (2001) 44:4216-4224
		GREGSON, S.J. et al., "Effect of C2/C3-endo unsaturation on the cytotoxicity and DNA-binding reactivity of pyrrolo-[2,1-c][1,4]-benzodiazepines," <i>Bioorg. Med. Chem. Lett.</i> (2000) 10(16):1849-1851
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		GREGSON, S.J. et al., "Synthesis of the first example of a C2-C3/C2'-C3'-endo unsaturated pyrrolo[2,1-c][1,4]benzodiazepine dimer," <i>Biorg. Med. Chem. Lett.</i> (2001) 11:2859-2862
		GREGSON, S.J. et al., "Synthesis of the first examples of A-C8/C-C2 amide-linked pyrrolo[2,1-c][1,4]benzodiazepine dimers," <i>Biorg. Med. Chem. Lett.</i> (2003) 13:2277-2280
		HAMBURGER, A.W. et al., "Primary bioassay of human tumor stem cells," <i>Science</i> (1977) 197:461-643

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		JAKOBSEN et al., "Design, synthesis, and pharmacological evaluation of thapsigargin analogues for targeting apoptosis to prostatic cancer cells," J. Med. Chem. (2001) 44:4696-4703			
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		LANGLOIS, N. et al., "Synthesis and cytotoxicity on sensitive and doxorubicin-resistant cell lines of new pyrrolo[2,1-c][1,4]benzodiazepines related to anthramycin," J. Med. Chem. (2001) 44:3754-3757			

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		LIPSHUTZ, B.H. et al., "Pd(II)_Catalyzed Acetal/Ehtal Hydrolysis/Exchange Reactions," Tetrahedron Lett. (1985) 26(6):705-708			
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.		MISCHIATI, C. et al., "Binding of hybrid molecules containing pyrrolo [2,1-c][1,4]benzodiazepine (PBD) and oligopyrrole carriers to the human immunodeficiency type 1 virus TAR-RNA," Biochem. Pharmacol. (2004) 67(3):401-410			
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		THURSTON, D.E., "Nucleic acid targeting: therapeutic strategies for the 21st century," Brit. J. Cancer (1999) 80(1):65-85			
.		TIBERGHIEN, A.C. et al., "Application of the stille coupling reaction to the synthesis of C2-substituted endo-exo unsaturated pyrrolo[2,1-c][1,4]benzodiazepines (PBDs)," Biorg. Med. Chem. Lett. (2004) 14:5041-5044			
.		WELLS, G. et al., "Pyrrolobenzodiazepine-polyamide libraries: synthesis and DNA binding selectivity," Proc. Am. Assoc. Canc. Res. (2003) 44:85-86, #452			
		WERMUTH et al., "Molecular Variations Based on Isosteric Replacements," The Practice of Medicinal Chemistry, Chapter 13 (1996) 203-237			
		WILLIAMS, M.A. et al., "Synthesis of conformationally constrained DTPA analogues. Incorporation of the ethylenediamine units as aminopyrrolidines," J. Org. Chem. (1994) 59(13):3616-3625			

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